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## **CLAIMS**

## 1. A compound of the formula

$$R_4X$$
 $Z$ 
 $R_6$ 
 $R_3X$ 
 $N$ 
 $XR_2$ 
 $R_6$ 
 $XR_1$ 
 $XR_2$ 

5 wherein:

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n is an integer of from 0 to 2;

 $Z \text{ is } N, N(O), O, S, S(O), S(O)_2, P, P(O), P(O)_2, Si, Si(O), or Si(O)_2;$ 

each X is independently C, C(O), N, N(O), O, S, S(O), S(O)<sub>2</sub>, P, P(O), P(O)<sub>2</sub>, Si, Si(O), or Si(O)<sub>2</sub> or is a bond; and

each of  $R_1$  to  $R_6$  is independently a bond or is selected from the group consisting of:

hydrogen;

halogen;

straight chain, cyclic, branched, substituted, heterocyclic, heteroatom substituted or unsubstituted alkyl, alkenyl, alkynyl, aryl, or heteroaryl;

phosphoryl groups such as phosphate, thiophosphate -O-P(S)(OH)<sub>2</sub>; phosphate esters -O-P(O)(OR)<sub>2</sub>; thiophosphate esters -O-P(S)(OR)<sub>2</sub>; phosphonate

-O-P(O)OHR; thiophosphonate -O-P(S)OHR; substituted phosphonate

-O-P(O)OR<sub>1</sub>R<sub>2</sub>; substituted thiophosphonate -O-P(S)OR<sub>1</sub>R<sub>2</sub>; -O-P(S)(OH)(SH); and cyclic phosphate;

other phosphorus containing compounds such as phosphoramidite

-O-P(OR)-NR<sub>1</sub>R<sub>2</sub>; and phosphoramidate -O-P(O)(OR)-NR<sub>1</sub>R<sub>2</sub>;

sulfur groups such as -O-S(O)(OH), -SH, -SR, -S( $\rightarrow$ O)-R, S(O)<sub>2</sub>R, RO-S(O)<sub>2</sub>,

-O-SO<sub>2</sub>NH<sub>2</sub>, -O-SO<sub>2</sub>R<sub>1</sub>R<sub>2</sub> or sulfamide –NHSO<sub>2</sub>NH<sub>2</sub>;

amino groups such as -NHR, -NR<sub>1</sub>R<sub>2</sub>, -NHAc, -NHCOR, -NH-O-COR, -

NHSO<sub>3</sub>, -NHSO<sub>2</sub>R, -N(SO<sub>2</sub>R)<sub>2</sub>, and/or amidino groups such as -NH-

C(=NH)NH<sub>2</sub> and/or ureido groups such as -NH-CO-NR<sub>1</sub>R<sub>2</sub> or thiouriedo groups such as -H-C(S)-NH<sub>2</sub>;

another unit of the structure I, attached through any position, where Z, X and  $R_1$  to  $R_6$  are as defined above; or

a substructure based upon a group of the following formula:

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wherein:

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Y is a bond or is selected from the group consisting of: straight chain, cyclic, branched, substituted, heterocyclic, heteroatom substituted or unsubstituted alkyl; straight chain, cyclic, branched, substituted, heterocyclic, heteroatom substituted or unsubstituted acyl; and aryl, substituted aryl, heteroaryl;

and

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each of  $R_7$  to  $R_{11}$  is independently at least one structure according to formula **I**, or a structure according to formula **II**;

with the provisos that:

when Z is O, and X is O or a bond, then all of  $R_1$  to  $R_5$  are not H or CH<sub>2</sub>OH; or when Z is N and X is O or a bond, then all of  $R_1$  to  $R_6$  are not H.

- 2. A compound according to claim 1, wherein said compound is PG2024, PG2037, PG2046, PG2155, as hereinbefore described.
  - 3. A compound according to claim 1, wherein said compound is any one of the compounds of Tables 1-4 of the description.
- 4. A pharmaceutical or veterinary composition for the prevention or treatment in a mammalian subject of a disorder resulting from angiogenesis, metastasis, inflammation, coagulation, thrombosis, and/or microbial infection, which composition comprises at least one compound according to claim 1 together with a pharmaceutically or veterinarially acceptable carrier or diluent for said at least one compound.
- 5. The composition according to claim 4 which further includes a pharmaceutically or veterinarially acceptable excipient, buffer, stabiliser, isotonicising agent, preservative or antioxidant.
  - 6. The composition according to claim 4, wherein said compound is present therein as an ester, a free acid or base, a hydrate, or a prodrug.
- 7. Use of a compound according to claim 1 in the manufacture of a medicament for the prevention or treatment in a mammalian subject of a disorder resulting from angiogenesis, metastasis, inflammation, coagulation, thrombosis, and/or microbial infection.

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- 8. The use according to claim 7, wherein said mammalian subject is a human subject.
- 9. A method for the prevention or treatment in a mammalian subject of a disorder resulting from angiogenesis, metastasis, inflammation, coagulation, thrombosis, and/or microbial infection, which method comprises administering to the subject an effective amount of at least one compound according to claim 1, or a composition comprising said at least one compound.

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- 10. The method according to claim 9 wherein said mammalian subject is a human subject.
- 11. The method according to claim 9, wherein said disorder resulting from angiogenesis is a proliferative retinopathy or angiogenesis resulting from the growth of a solid tumour.
- 10 12. The method according to claim 9, wherein said disorder resulting from inflammation is rheumatoid arthritis, multiple sclerosis, inflammatory bowel disease, allograft rejection or chronic asthma.
  - 13. The method according to claim 9, wherein said disorder resulting from coagulation and/or thrombosis is deep venous thrombosis, pulmonary embolism, thrombotic stroke, peripheral arterial thrombosis, unstable angina or myocardial infarction.
  - 14. The method according to claim 9, wherein said disorder resulting from viral infection is Herpes Simplex.